

APPLICANT FACSIMILE OF FORM PTO-1449
REV 7-80U.S. DEPARTMENT OF COMMERCE
PATENT AND TRADEMARK OFFICE

ATTY DOCKET NO

IGI-001CN3

SERIAL NO.

09/335,956

LIST OF PUBLICATIONS CITED BY APPLICANT
(Use several sheets if necessary)

APPLICANT

David C. Ward et al.

FILING DATE

06-18-99

GROUP

U.S. PATENT DOCUMENTS

EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
M	AA	4,358,535	11/82	Falkow et al.	435	5	
M	AB	4,647,529	03/87	Rodland et al.	435	6	
M	AC	4,681,840	07/87	Stephenson et al.	435	6	
M	AD	4,683,195	07/87	Mullis et al.	435	6	
M	AE	4,707,440	11/87	Stavrianopoulos	435	6	
M	AF	4,710,465	12/87	Weissman et al.	435	91	
M	AG	4,711,955	12/87	Ward et al.	536	29	
M	AH	4,721,669	01/88	Barton	435	6	
M	AI	4,725,536	02/88	Fritsch et al.	435	6	
M	AJ	4,755,458	07/88	Rabbani et al.	435	5	
M	AK	4,770,992	09/88	Van den Engh et al.	435	6	
M	AL	4,772,691	09/88	Herman	536	27	
M	AM	4,888,278	12/89	Singer et al.	435	6	

FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	TRANSLATION YES NO
M	AN	GB 2 019 408	10/79	U.K.			
M	AO	WO 87/05027	08/87	PCT			
M	AP	GB 2 215 724	09/89	U.K.			

OTHERS (including Author, Title, Date, Pertinent Pages, Etc.)

M	AQ	Albertson, D., "Localization of the Ribosomal Genes in <i>Caenorhabditis Elegans</i> Chromosomes by <i>in Situ</i> Hybridization Using Biotin-Labeled Probes," <i>EMBO Journal</i> , 3(6):1227-1234 (1984);
M	AR	Albertson, D., "Mapping Muscle Protein Genes by <i>in Situ</i> Hybridization Using Biotin-Labeled Probes," <i>EMBO Journal</i> , 4(10):2493-2498 (1985);
M	AS	Ardeshir, F., et al., "Structure of Amplified DNA in Different Syrian Hamster Cell Lines Resistant to N-(Phosphonacetyl)-L-Aspartate," <i>Molecular and Cellular Biology</i> , 3(11):2076-2088 (1983);
M	AT	Arnoldus, E.P.J., et al., "Detection of the Philadelphia Chromosome in Interphase Nuclei (With 2 Color Plates)," <i>Cytogenet Cell Genet</i> , 54:108-111 (1990);
M	AU	Bar-Am, I., et al., "Detection of Amplified DNA Sequences in Human Tumor Cell Lines by Fluorescence <i>in Situ</i> Hybridization," <i>Genes, Chromosomes & Cancer</i> , 4:314-320 (1992);

Examiner

M

Date Considered

22 May 1990

*EXAMINER

Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

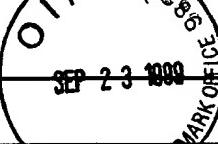
APPLICANT FACSIMILE OF FORM PTO-1449
REV 7-80U.S. DEPARTMENT OF COMMERCE
PATENT AND TRADEMARK OFFICE

ATTY DOCKET NO

IGI-001CN3

SERIAL NO.

09/335,956

LIST OF PUBLICATIONS CITED BY APPLICANT
(Use several sheets if necessary)

APPLICANTS

David C. Ward et al.

FILING DATE

06-18-99

GROUP

OTHERS (including Author, Title, Date, Pertinent Pages, Etc.)

<i>PATENT</i>	<i>BA</i>	Benton, W. and Davis, R., "Screening for Recombinant Clones by Hybridization to Single Plaques in situ," <i>Science</i> , 196:180-182 (1977);
<i>M</i>	<i>BB</i>	Bergerheim, U., et al., "Deletion Mapping in Human Renal Cell Carcinoma ¹ ," <i>Cancer Research</i> , 49:1390-1396 (1989);
<i>M</i>	<i>BC</i>	Bookstein, R., et al., "Human Retinoblastoma Susceptibility Gene: Genomic Organization and Analysis of Heterozygous Intragenic Deletion Mutants," <i>PNAS (USA)</i> , 85:2210-2214 (1988);
<i>M</i>	<i>BD</i>	Brison, O., et al., "General Method for Cloning Amplified DNA by Differential Screening with Genomic Probes," <i>Molecular and Cellular Biology</i> , 2(15):578-587 (1982);
<i>M</i>	<i>BE</i>	Britten, R., et al., "Analysis of Repeating DNA Sequences by Reassociation" <i>Methods of Enzymology</i> , 29:363-418 (1974);
<i>M</i>	<i>BF</i>	Buongiorno-Nardelli, M., "Autoradiographic Detection of Molecular Hybrids between rRNA and DNA in Tissue Sections," <i>Nature</i> , 225:946-948 (1970);
<i>M</i>	<i>BG</i>	Cannizzaro, L.A., et al., "In Situ Hybridization and Translocation Breakpoint Mapping," <i>Cytogenet Cell Genet</i> , 39:173-178 (1985);
<i>M</i>	<i>BH</i>	Cohen, A., et al., "Hereditary Renal-Cell Carcinoma Associated with a Chromosomal Translocation," <i>The New England Journal of Medicine</i> , 301(11):592-595 (1979);
<i>M</i>	<i>BI</i>	Collins, F. and Weissman, S., "Directional Cloning of DNA Fragments at a Large Distance from an Initial Probe: A Circularization Method," <i>PNAS (USA)</i> , 81:6812-6816 (1984);
<i>M</i>	<i>BJ</i>	Cox, K., et al., "Detection of mRNAs in Sea Urchin Embryos by <i>in Situ</i> Hybridization Using Asymmetric RNA Probes," <i>Developmental Biology</i> , 101:485-502 (1984);
<i>P</i>	<i>BK</i>	Cremer, T., et al., "Detection of Chromosome Aberrations in the Human Interphase Nucleus by Visualization of Specific Target DNAs with Radioactive and Non-Radioactive <i>In Situ</i> Hybridization Techniques: Diagnosis of Trisomy 18 with Probe L1.84," <i>Hum. Genet.</i> , 74:346-352 (1986);
<i>M</i>	<i>BL</i>	Cremer, T., et al., "Rapid Interphase and Metaphase Assessment of Specific Chromosomal Changes in Neuroectodermal Tumor Cells by <i>in Situ</i> Hybridization with Chemically Modified DNA Probes," <i>Experimental Cell Research</i> , 176:199-220 (1988);
<i>M</i>	<i>BM</i>	Cremer, T., et al., "Rapid Metaphase and Interphase Detection of Radiation-Induced Chromosome Aberrations in Human Lymphocytes by Chromosomal Suppression <i>In Situ</i> Hybridization," <i>Cytometry</i> , 11:110-118 (1990);
<i>M</i>	<i>BN</i>	Devilee, P., et al., "Detection of Chromosome Aneuploidy in Interphase Nuclei from Human Primary Breast Tumors Using Chromosome-specific Repetitive DNA Probes," <i>Cancer Research</i> , 48:5825-5830 (1988);
<i>M</i>	<i>BO</i>	Durnam, D., et al., "Detection of Species Specific Chromosomes in Somatic Cell Hybrids," <i>Som. Cell Molec. Genetics</i> , 11(6):571-577 (1985);
<i>M</i>	<i>BP</i>	Erikson, J., et al., "Heterogeneity of Chromosome 22 Breakpoint in Philadelphia-Positive(Ph+) Acute Lymphocytic leukemia," <i>PNAS USA</i> , 83:1807-1811 (1986);
<i>M</i>	<i>BQ</i>	Fisher, J.H., et al., "Molecular Hybridization Under Conditions of High Stringency Permits Cloned DNA Segments Containing Reiterated DNA Sequences to be Assigned to Specific Chromosomal Locations," <i>PNAS USA</i> , 81:520-524 (1984);
<i>M</i>	<i>BR</i>	Friend, S., et al., "A Human DNA Segment with Properties of the Gene that Predisposes to Retinoblastoma and Osteosarcoma," <i>Nature</i> , 323:643-646 (1986);

Examiner

Date Considered

27 Meier

*EXAMINER

Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

LIST OF PUBLICATIONS CITED BY APPLICANT
(Use several sheets if necessary)

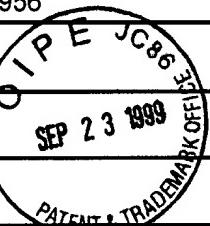
APPLICANTS

David C. Ward et al.

FILING DATE

06-18-99

GROUP



OTHERS (including Author, Title, Date, Pertinent Pages, Etc.)

<i>M</i>	CA	Fuscoe, J., et al., "An Efficient Method for Selecting Unique-Sequence Clones from DNA Libraries and Its Application to Fluorescent Staining of Human Chromosome 21 Using <i>in Situ</i> Hybridization," <i>Genomics</i> , 5:100-109 (1989);
<i>N</i>	CB	Gall, J. and Pardue, M., "Formation and Detection of RNA -DNA Hybrid Molecules in Cytological Preparations," <i>PNAS USA</i> , 63:378-383 (1969);
<i>M</i>	CC	Gray, J.W., et al., "Fluorescence Hybridization to Human Chromosome 21 Using Probes from a Charon 21 A Library," <i>Cytometry</i> , (Suppl. 1), Abst. 19, p. 4 (1987);
<i>M</i>	CD	Gray, J.W., et al., "Quantitative Cytogenetics: Progress Report on the Development of Fluorescence Hybridization for Specific Chromosome Staining," (Abstract) Lawrence Livermore National Laboratory, Livermore, CA (UCRL 93567);
<i>M</i>	CE	Grunstein, M., et al., "Colony Hybridization: A Method for the Isolation of Cloned DNAs That Contain a Specific Gene," <i>PNAS, USA</i> , 72(10):3961-3965 (1975);
<i>M</i>	CF	Harper, M. and Saunders, G., "Localization of Single Copy DNA Sequences on G-Banded Human Chromosomes by <i>in Situ</i> Hybridization," <i>Chromosoma (Berl.)</i> , 83:431-439 (1981);
<i>M</i>	CG	Harper, M., et al., "Localization of the Human Insulin Gene to the Distal end of the Short Arm of Chromosome 11," <i>PNAS USA</i> , 78(7):4458-4460 (1981);
<i>M</i>	CH	Herzenberg, L., et al., "Fetal Cells in the Blood of Pregnant Women: Detection and Enrichment by Fluorescence-Activated Cell Sorting," <i>PNAS USA</i> , 76(3):1453-1455 (1979);
<i>M</i>	CI	Hood, L., et al., <i>Molecular Biology of Eucaryotic Cells</i> , W.A. Benjamin, Inc. Menlo Park, Calif., CH. 2-10, 47-51 (1975);
<i>M</i>	CJ	Jabs, E., et al., "Characterization of a Cloned DNA Sequence That is Present at Centromeres of all Human Autosomes and the X Chromosome and Shows Polymorphic Variation," <i>PNAS USA</i> , 81:4884-4888 (1984);
<i>M</i>	CK	John, H.A., et al., "RNA-DNA Hybrids at the Cytological Level," <i>Nature</i> , 223:582-587 (1969);
<i>M</i>	CL	Kao, F., et al., "Assignment of the Structural Gene Coding for Albumin to Human Chromosome 4," <i>Human Genetics</i> , 62:337-341 (1982);
<i>M</i>	CM	Kievits, T., et al., "Direct Nonradioactive <i>in Situ</i> Hybridization of Somatic Cell Hybrid DNA to Human Lymphocyte Chromosomes," <i>Cytometry</i> , 11:105-109 (1990);
<i>M</i>	CN	Landegent, J.E., et al., "2 -Acetylaminofluorene-modified Probes for the Indirect Hybridocytochemical Detection of Specific Nucleic Acid Sequences," <i>Experimental Cell Research</i> , 153:61-72 (1984);
<i>M</i>	CO	Landegent, J.E., et al., "Chromosomal Localization of a Unique Gene by Non-Autoradiographic <i>in Situ</i> Hybridization," <i>Nature</i> , 317:175-177 (1985);
<i>M</i>	CP	Landegent, J.E., et al., "Use of Whole Cosmid Cloned Genomic Sequences for Chromosomal Localization by Non-Radioactive <i>in Situ</i> Hybridization," <i>Human Genetics</i> , 77:366-370 (1987);
<i>M</i>	CQ	Langer-Safer, P., et al., "Immunological Method for Mapping Genes on <i>Drosophila</i> Polytene Chromosomes," <i>PNAS USA</i> , 79:4381-4385 (1982);
<i>M</i>	CR	Lawrence, J., et al., "Sensitive, High-Resolution Chromatin and Chromosome Mapping <i>In Situ</i> : Presence and Orientation of Two Closely Integrated Copies of EBV in a Lymphoma Line," <i>Cell</i> , 52:51-61 (1988);
Examiner		Date Considered <i>22 May 2002</i>
*EXAMINER		Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

APPLICANT FACSIMILE OF FORM PTO-1449 REV 7-80	U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTY DOCKET NO IGI-001CN3	SERIAL NO. 09/335,956
LIST OF PUBLICATIONS CITED BY APPLICANT (Use several sheets if necessary)		APPLICANTS David C. Ward et al.	O P E JC 86 SEP 23 1999 PATENT & TRADEMARK OFFICE
		FILING DATE 06-18-99	

OTHERS (including Author, Title, Date, Pertinent Pages, Etc.)

<i>p</i>	DA	Lawrence Livermore National Laboratory, "Chromosome- Specific Human Gene Libraries," <i>Energy and Technology Review</i> , 82-83 (1985);
<i>dl</i>	DB	Lawrence Livermore National Laboratory, "Fluorescent Labeling of Human Chromosomes with Recombinant DNA Probes," <i>Energy and Technology Review</i> (UCRL 5200-85-7), 84-85 (1985);
<i>ll</i>	DC	LeGrys, V., et al., "Clinical Applications of DNA Probes in the Diagnosis of Genetic Diseases," <i>CRC Critical Reviews in Clinical Laboratory Sciences</i> , 25(4):255-274 (1987);
<i>m</i>	DD	Lewin, B. (editor), <i>Eukaryotic Genomes: A Continuum of Sequences</i> (Chapter 18), <i>Genes</i> (2nd Edition John Wiley & Sons, Inc.), 293-298 and 464-465 (1984);
<i>r</i>	DE	Lewin, R., "Genetic Probes Become Ever Sharper" <i>Science</i> , 221(4616):1167 (1983);
<i>r</i>	DF	Lichter, P., et al., "High-Resolution Mapping of Human Chromosome 11 by in Situ Hybridization with Cosmid Clones," <i>Science</i> , 247:64-69 (1990);
<i>r</i>	DG	Lichter, P., et al., "Is Non-isotopic <i>in Situ</i> Hybridization Finally Coming of Age?," <i>Nature</i> , 345:93-94 (1990);
<i>o</i>	DH	Lichter, P., et al., "Rapid Detection of Human Chromosome 21 Aberrations by <i>in Situ</i> Hybridization," <i>PNAS USA</i> , 85:9664-9668 (1988);
<i>m</i>	DI	Litt, M. and White, R.L., "A Highly Polymorphic Locus in Human DNA Revealed by Cosmid-Derived Probes," <i>PNAS USA</i> , 82:6206-6210 (1985);
<i>m</i>	DJ	Manuelidis, L. and Ward, D., "Chromosomal and Nuclear Distribution of the HindIII 1.9-kb Human DNA Repeat Segment," <i>Chromosoma (Berl.)</i> , 91:28-38 (1984);
<i>m</i>	DK	Manuelidis, L., "Individual Interphase Chromosome Domains Revealed by <i>in Situ</i> Hybridization" <i>Hum. Genet.</i> , 71:288-293 (1985);
<i>m</i>	DL	Manuelidis, L., "Different Central Nervous System Cell Types Display Distinct and Nonrandom Arrangements of Satellite DNA Sequences" <i>PNAS USA</i> , 81:3123-3127 (1984);
<i>m</i>	DM	McCormick, F., "The Polymerase Chain Reaction and Cancer Diagnosis," <i>Cancer Cells</i> , 1(2), 56-61 (1989);
<i>o</i>	DN	Montgomery, K., et al., "Specific DNA Sequence Amplification in Human Neuroblastoma Cells," <i>PNAS USA</i> , 80:5724-5728 (1983);
<i>m</i>	DO	Nederlof, P., et al., "Detection of Chromosome Aberrations in Interphase Tumor Nuclei by Nonradioactive <i>In Situ</i> Hybridization," <i>Cancer Genet Cytogenet</i> , 42:87-98 (1989);
<i>m</i>	DP	Olsen, A., et al., "Isolation of Unique Sequence Human X Chromosomal Deoxyribonucleic Acid," <i>Biochemistry</i> , 19:2419-2428 (1980);
<i>m</i>	DQ	Pinkel, D., et al., "Cytogenetic Analysis by <i>in Situ</i> Hybridization With Fluorescently Labeled Nucleic Acid Probes," <i>Cold Spring Harbor Symposia on Quantitative Biology</i> , L1:151-157 (1986);
<i>m</i>	DR	Pinkel, D., et al., "Cytogenetic Analysis During Leukemia Therapy Using Fluorescence <i>in Situ</i> Hybridization with Chromosome-Specific Nucleic Acid Probes," <i>Am. J. Hum. Genet.</i> (Supplement), 41(3), A34 (096; 12.12) (1987);
<i>p</i>	DS	Pinkel, D., et al., "Cytogenetic Analysis Using Quantitative, High-Sensitivity, Fluorescence Hybridization," <i>PNAS USA</i> , 83:2934-2938 (1986);
<i>r</i>	DT	Pinkel, D. et al., "Cytogenetics Using Fluorescent Nucleic Acid Probes and Quantitative Microscopic Measurement," (UCRL 93269 Abstract), <i>Analytical Cytology X Conference</i> , Hilton Head Resort, SC, (November 17-22, 1985);

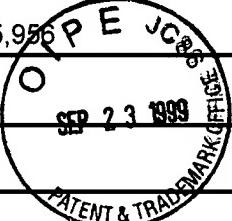
Examiner

Date Considered

22 May 2000

*EXAMINER

Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

LIST OF PUBLICATIONS CITED BY APPLICANT
(Use several sheets if necessary)

APPLICANTS

David C. Ward et al.

FILING DATE

06-18-99

GROUP

OTHERS (including Author, Title, Date, Pertinent Pages, Etc.)

P	EA	Pinkel, D., et al., "Detection of Structural and Numerical Abnormalities in Metaphase Spreads and Interphase Nuclei Using in Situ Hybridization," <i>Cancer Genet and Cytogenet</i> (UCRL 101043 Abstract 34), 41:236 (1989);
V	EB	Pinkel, D., et al., "Detection of Structural Chromosome Aberrations in Metaphase Spreads and Interphase Nuclei by in Situ Hybridization High Complexity Probes Which Stain Entire Human Chromosomes," <i>Am. J. Hum. Genet.</i> (Supplement), 43:3, A118 (Abstract (0471)11.5) (September 1988);
B	EC	Pinkel, D., et al., "Fluorescence in Situ Hybridization with Human Chromosome-Specific Libraries: Detection of Trisomy 21 and Translocations of Chromosome 4," <i>PNAS USA</i> , 85:9138-9142 (1988);
J	ED	Pinkel, D., et al., "Genetic Analysis by Quantitative Microscopy and Flow Cytometry Using Fluorescence in Situ Hybridization with Chromosome-Specific Nucleic Acid Probes," <i>Am. J. Hum. Genet.</i> (Supplement), Vol 39:3, A129 (379) (Sept. 1986);
N	EE	Pinkel, D., et al., "Rapid, Quantitative Cytogenetic Analysis Using Fluorescently Labeled Nucleic Acid Probes," (UCRL 93553 Abstract), U.S. - Japan Joint Environmental Panel Conf., Research Triangle Park, NC, (October 21-23, 1985);
V	EF	Pinkel, D., et al., "Simplified Cytogenetics Using Biotin Labeled Nucleic Acid Probes and Quantitative Fluorescence Microscopy," (UCRL 93243 Abstract), <i>American Journal of Human Genetics</i> (Supplement), 37(4):A112 (July 1985);
M	EG	Rappold, G.A., et al., "Sex Chromosome Positions in Human Interphase Nuclei as Studied by in Situ Hybridization with Chromosome Specific DNA Probes," <i>Human Genetics</i> , 67:317-325 (1984);
M	EH	Roelofs, H., et al., "Gene Amplification in Human Cells May Involve Interchromosomal Transposition and Persistence of the Original DNA Region," <i>The New Biologist</i> , 4(1):75-86 (1992);
M	EI	Scalenghe, F., et al., "Microdissection and Cloning of DNA from a Specific Region of <i>Drosophila melanogaster</i> Polytene Chromosomes," <i>Chromosoma (Berl.)</i> , 82:205-216 (1981);
M	EJ	Schardin, M., et al., "Specific Staining of Human Chromosomes in Chinese Hamster X man Hybrid Cell Lines Demonstrates Interphase Chromosome Territories," <i>Hum Genet</i> , 71:281-287 (1985);
M	EK	Schmeckpeper, b., et al., "Partial purification and characterization of DNA from the human X chromosome," <i>PNAS USA</i> , 76(12):6525-6528 (1979);
V	EL	Sealy et al., "Removal of repeated sequences from hybridisation probes," <i>Nucleic Acids Research</i> , 13(6):1905-1922 (1985);
M	EM	Selypes et al., "A Noninvasive Method for Determination of the Sex and Karyotype of the Fetus From the Maternal Blood," <i>Hum. Genet.</i> , 79:357-359 (1988);
M	EN	Smith et al., "Distinctive Chromosomal Structures Are Formed Very Early in the Amplification of CAD Genes in Syrian Hamster Cells," <i>Cell</i> , 63:1219-1227 (1990);
V	EO	Sparks et al., "Regional Assignment of Genes for Human Esterase D and Retinoblastoma in Chromosome Band 13q14," <i>Science</i> , 208:1042-1044 (1988);

Examiner

Date Considered

*EXAMINER

Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

LIST OF PUBLICATIONS CITED BY APPLICANT
(Use several sheets if necessary)

APPLICANTS

David C. Ward et al.

FILING DATE

06-18-99

GROUP

SEP 23 1999

P A T E N T & T R A D E M A R K O F F I C E

OTHERS (including Author, Title, Date, Pertinent Pages, Etc.)

<i>✓</i>	FA	Stewart et al., "Cloned DNA probes regionally mapped to human chromosome 21 and their use in determining the original of nondisjunction," <i>Nucleic Acids Research</i> , 13(11):4125-4132 (1985);
<i>✓</i>	FB	Straume et al., "Chromosome translocations at low radiation doses quantified using fluorescent DNA probes," Radiation Research Society Meeting (UCRL 93837 Abstract), Las Vegas, NV (April 12-17, 1986);
<i>✓</i>	FC	Szabo and Ward, "Emerging Techniques. What's new with hybridization in situ?" <i>TIBS</i> , 7(11):425-427 (1982);
<i>✓</i>	FD	Trask et al., "Detection of DNA sequences in nuclei in suspension by in situ hybridization and dual beam flow cytometry," Analytical Cytology X Conference (UCRL 93372 Abstract), Hilton Head Resort, SC (November 17-22, 1985);
<i>✓</i>	FE	Trask et al., "The Proximity of DNA Sequences in Interphase Cell Nuclei is Correlated to Genomic Distance and Permits Ordering of Cosmids Spanning 250 Kilobase Pairs," <i>Genomics</i> , 5:710-717 (1989);
<i>✓</i>	FF	Trask et al., "Early dihydrofolate reductase gene amplification events in CHO cells usually occur on the same chromosome arm as the original locus," <i>Genes & Development</i> , 3:1913-1925 (1989);
<i>✓</i>	FG	Trent et al., "Report of the committee on structural chromosome changes in neoplasia," <i>Cytogenet Cell Genet</i> , 51:533-562 (1989);
<i>✓</i>	FH	Van Dilla and Deaven (Abstract), "Construction and Availability of Human Chromosome-Specific DNA Libraries from Flow Sorted Chromosomes: Status Report," <i>Am J. of Human Genetics</i> , 37, (R. Supplement) (July 1985);
<i>✓</i>	FI	Wallace et al., "The use of synthetic oligonucleotides as hybridization probes. II. Hybridization of oligonucleotides of mixed sequence to rabbit β-globin DNA," <i>Nucleic Acids Research</i> , 9(4):879-894 (1981);
<i>✓</i>	FJ	Weiss et al., "Organization and evolution of the class I gene family in the major histocompatibility complex of the C57BL/10 mouse," <i>Nature</i> , 310:650-655 (1984);
<i>✓</i>	FK	Willard et al., "Isolation and characterization of a major tandem repeat family from the human X chromosome," <i>Nucleic Acids Research</i> , 11(7):2017-2033 (1983);
<i>✓</i>	FL	Windle et al., "A central role for chromosome breakage in gene amplification, deletion formation, and amplicon integration" <i>Genes & Development</i> , 5:160-174 (1991);
<i>✓</i>	FM	Yunis et al., "Localization of Sequences Specifying Messenger RNA to Light-Staining G-Bands of Human Chromosomes" <i>Chromosoma (Berl.)</i> , 61:335-344 (1977).
	FN	
	FO	
	FP	
	FQ	
	FR	

Examiner

Date Considered

22 May 2000

*EXAMINER

Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.